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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO	
09/682,443	09/04/2001	Michiel Jacques van Nieuwstadt	200-1758 JDR	9487	
22844	7590 12/28/2004		EXAMINER		
FORD GLOBAL TECHNOLOGIES, LLC.			NGUYEN, TU MINH		
SUITE 600 - ONE PARKL	PARKLANE TOWERS ANE BLVD	EAST	ART UNIT	PAPER NUMBER	
DEARBORN,		-	3748		

DATE MAILED: 12/28/2004

Please find below and/or attached an Office communication concerning this application or proceeding.

Office Action Summary		Application No. Applicant(s)						
		09/682,443	VAN NIEUWSTAI JACQUES	VAN NIEUWSTADT, MICHIEL JACQUES				
		Examiner	Art Unit					
		Tu M. Nguyen	3748					
The MAILING DATE of this Period for Reply	s communication app	ears on the cover sheet wi	th the correspondence ac	idress				
A SHORTENED STATUTORY F THE MAILING DATE OF THIS C - Extensions of time may be available under to after SIX (6) MONTHS from the mailing data. - If the period for reply specified above, the fixed particular to reply within the set or extended power particular to reply within the set or extended power particular to reply within the set or extended power particular to reply within the set or extended power particular to reply within the set or extended power particular to reply within the set or extended power particular to reply within the set or extended power particular to reply within the set or extended power particular to reply within the set or extended power particular to reply within the set or extended power particular to reply within the set or extended power particular to reply within the set or extended power particular to reply within the set or extended power particular to reply within the set or extended power particular to reply set or particular to reply set ore	COMMUNICATION. the provisions of 37 CFR 1.13 e of this communication. s than thirty (30) days, a reply e maximum statutory period w eriod for reply will, by statute, hree months after the mailing	6(a). In no event, however, may a rowithin the statutory minimum of thirtill apply and will expire SIX (6) MON cause the application to become AB	eply be timely filed y (30) days will be considered time THS from the mailing date of this c ANDONED (35 U.S.C. § 133).					
Status								
1) Responsive to communica	tion(s) filed on 17 Au	gust_2004.						
2a)⊠ This action is FINAL .		action is non-final.						
3)☐ Since this application is in	condition for allowan	ce except for formal matte	ers, prosecution as to the	e merits is				
closed in accordance with	closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.							
Disposition of Claims								
4)⊠ Claim(s) 1 and 4-21 is/are	Claim(s) <u>1 and 4-21</u> is/are pending in the application.							
	4a) Of the above claim(s) is/are withdrawn from consideration.							
5) Claim(s) is/are allow	Claim(s) is/are allowed.							
6)⊠ Claim(s) <u>1 and 4-21</u> is/are	Claim(s) 1 and 4-21 is/are rejected.							
7) Claim(s) is/are obje								
8) Claim(s) are subject	t to restriction and/or	election requirement.						
Application Papers								
9)⊠ The specification is objecte	d to by the Examiner	:						
10)⊠ The drawing(s) filed on <u>24 September 2003</u> is/are: a)⊠ accepted or b)□ objected to by the Examiner.								
· · · · · · · · · · · · · · · · · · ·	Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a). Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).							
i i i i i i i i i i i i i i i i i i i								
11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.								
Priority under 35 U.S.C. § 119								
12) Acknowledgment is made of	of a claim for foreign	oriority under 35 U.S.C. &	. 119(a)-(d) or (f)					
a) ☐ All b) ☐ Some * c) ☐ N		priority dilder 55 C.C.C. S	110(a)-(u) or (i).					
·- <u>-</u> ·-								
	•	have been received in A	nnlication No					
	•	ty documents have been	• • • • • • • • • • • • • • • • • • • •	Stage				
·	International Bureau			- 90				
* See the attached detailed O			received.					
Attachment(s)		م السيد	(BTO 443)					
 Notice of References Cited (PTO-892) Notice of Draftsperson's Patent Drawin 	g Review (PTO-948)		Summary (PTO-413) s)/Mail Date					
3) Information Disclosure Statement(s) (P Paper No(s)/Mail Date		5) ☐ Notice of Ir 6) ☐ Other:	nformal Patent Application (PT 	O-152)				

Application/Control Number: 09/682,443 Page 2

Art Unit: 3748

DETAILED ACTION

1. An Applicant's Amendment filed on August 7, 2004 has been entered. Claims 20-21 have been added. Overall, claims 1 and 4-21 are pending in this application.

Drawings

2. The formal drawing of Figure 2 filed on September 24, 2003 has been approved for entry.

Specification

3. The disclosure is objected to because on page 4, paragraph 0014, the sentence is incomplete. According to PTO's record, paragraph 0014 reads "FIG. 2 is a functional block diagram of an engine exhaust system according to the", which is clearly an incomplete sentence. Appropriate correction is required.

Claim Objections

- 4. Claim 21 is objected to because
 - On line 7 of the claim, "thetemperature" should read -- the temperature--.
 - On the last line of the claim, "again" should read --aging--.

Appropriate correction is required.

Art Unit: 3748

Claim Rejections - 35 USC § 102

5. The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office Action:

A person shall be entitled to a patent unless -

- (b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.
- 6. Claims 1, 4, 12, and 20 are rejected under 35 U.S.C. 102(b) as being anticipated by Bagley et al. (U.S. Patent 5,497,617).

Re claim 1, as illustrated in Figure 2 and indicated in claims 1-2, Bagley et al. disclose a method for controlling hydrocarbon (methanol) injection into an engine exhaust to reduce NOx, comprising injecting the hydrocarbon into the engine exhaust in accordance with detection of a light-off event, such light-off event being detected when there is a hydrocarbon-oxygen reaction wherein an exothermic reaction is produced and detected (Figure 2 shows that a mixture of methanol and air has an ignition temperature at a catalyst temperature as low as 50°C and thus, exhibits an exotherm as shown; as claimed in claims 1-2, Bagley et al. utilize a heater to raise a catalyst temperature to an ignition temperature of a mixture of methanol and air, they then supply the mixture of methanol and air into the catalyst, resulting in the combustion of said mixture and raising the catalyst temperature until the catalyst reaches light-off temperatures for other components (NOx, HC, CO) in the exhaust gas).

Re claims 4, 12, and 20, as illustrated in Figure 2 and indicated in claims 1-2, Bagley et al. disclose a method for controlling hydrocarbon (methanol) injection into an engine exhaust to

Art Unit: 3748

reduce NOx in such exhaust, such engine exhaust with the NOx and the injected hydrocarbon being directed to a catalyst for reaction therein, comprising:

- (a) detecting an exothermic reaction across the catalyst (step b) in claim 1; also see Figure 2 where an exotherm for methanol is detected when a catalyst temperature reaches approximately 50°C);
- (b) detecting, measuring, or obtaining a temperature of the catalyst in response to the detected exothermic reaction (from Figure 2, a catalyst temperature of about 45°C is determined when an exotherm for methanol is detected); and
- (c) injecting the hydrocarbon into the reaction in accordance with the measured temperature (step c) in claim 1).
- 7. Claims 4-21 are rejected under 35 U.S.C. 102(b) as being anticipated by Hirota et al. (U.S. Patent 5,201,802).

Re claims 4, 12, 14, and 20, as shown in Figures 6 and 14, Hirota et al. disclose a method for controlling hydrocarbon injection into an engine exhaust to reduce NOx in such exhaust, such engine exhaust with the NOx and the injected hydrocarbon being directed to a catalyst (6) for reaction therein, comprising:

- (a) detecting an exothermic reaction across the catalyst (step 608);
- (b) detecting, measuring, or obtaining a temperature of an inlet of the catalyst in response to the detected exothermic reaction (step 608) (an inlet temperature t1 is detected and measured using an upstream temperature sensor (24)), and
- (c) injecting the hydrocarbon into the reaction in accordance with the measured temperature (steps 618 and 620).

Art Unit: 3748

Re claims 5, 6, 10, 13, 15, 16, and 19, as illustrated in Figures 6 and 14-18, Hirota et al. disclose a method for controlling hydrocarbon injection into an engine exhaust to reduce NOx in such exhaust, such engine exhaust with the NOx and the injected hydrocarbon being directed to a catalyst (6) for reaction therein, comprising:

Page 5

- (a) detecting a pair (t1 and t2) of temperatures across the catalyst to provide a temperature difference (Δt) indicating an exothermic reaction across the catalyst (step 608);
- (b) comparing the temperature difference with a predetermined temperature threshold (ΔTi) (step 610);
- (c) determining an exothermic condition temperature (T1) when the temperature difference is determined to exceed the threshold (step 614, Figure 17), such exothermic condition temperature being determined from an upstream one of the detected temperatures;
- (d) comparing the determined exothermic condition temperature with an exothermic condition temperature (550 in Figure 17) expected from the catalyst at a time prior to the determined exothermic condition temperature; and
- (e) modifying the injected hydrocarbon in accordance with the last-mentioned comparison (steps 618 and 620; also see Figure 18 and line 10 of column 9 to line 3 of column 10) (Hirota et al. determine in advance a desired lower limit catalyst inlet temperature T1 and a desired upper limit catalyst outlet temperature T2 for the optimum reduction of NOx as a function of the degradation extent DR (Figure 17). For a non-deteriorated catalyst, T1 and T2 equal 450 and 550, respectively. If a detected temperature difference (Δt) across the catalyst is different from a predetermined temperature threshold (ΔTi), a degradation extent DR is

Art Unit: 3748

calculated (step 612); and a set of desired temperature values T1 and T2 are determined based on the calculated DR (step 614). A hydrocarbon concentration H1 is also determined based on DR).

Re claim 21, as shown in Figures 6 and 14-18, Hirota et al. disclose a method for controlling hydrocarbon injection into an engine exhaust to reduce NOx in such exhaust, such engine exhaust with the NOx and the injected hydrocarbon being directed to a catalyst for reaction therein, comprising:

- (a) identifying catalyst light-off by detecting production of an exothermic reaction across the catalyst when a temperature difference (Δt) across the catalyst exceeds a threshold value (ΔTi) (steps 608 and 610);
- (b) determining a light-off temperature (T1) of the catalyst by measuring the temperature at which the exothermic reaction is detected (step 614, Figure 17);
- (c) obtaining a measure (DR) of catalyst aging based on the detected temperature (step 612), and
- (d) adjusting injection of the hydrocarbon into the reaction in accordance with the measure of catalyst aging (steps 618 and 620; also see Figure 18 and line 10 of column 9 to line 3 of column 10) (Hirota et al. determine in advance a desired lower limit catalyst inlet temperature T1 and a desired upper limit catalyst outlet temperature T2 for the optimum reduction of NOx as a function of the degradation extent DR (Figure 17). For a non-deteriorated catalyst, T1 and T2 equal 450 and 550, respectively. If a detected temperature difference (Δt) across the catalyst is different from a predetermined temperature threshold (ΔTi), a degradation extent DR is calculated (step 612); and a set of desired temperature values T1 and T2 are

Art Unit: 3748

determined based on the calculated DR (step 614). A hydrocarbon concentration H1 is also determined based on DR).

Re claims 7, 9, 17, and 18, as shown in Figures 6 and 14-18, Hirota et al. disclose a system and a processor (10) for controlling hydrocarbon injection into an engine exhaust to reduce NOx in such exhaust, such engine exhaust with the NOx and the injected hydrocarbon being directed to a catalyst (6) for reaction therein, the system comprising:

- (a) a catalyst (6) for facilitating a reaction between the injected hydrocarbon and NOx in the exhaust;
- (b) a hydrocarbon injector (14) for injecting the hydrocarbon into the exhaust upstream of the catalyst;
 - (c) a detecting system comprising:
- a pair of sensors (24, 20) each detecting a common parameter in the exhaust, one of such sensors being upstream of the catalyst and the other one of the sensors being downstream of the first sensor, and
- a processor (10) for controlling the hydrocarbon injector in response to the pair of sensors, such processor being programmed to:
- comparing a difference (Δt) in the common parameter detected by the pair of sensors with a predetermined temperature threshold (ΔTi) (step 610);
- determining an exothermic condition temperature (T1) from an upstream sensor (24) when the difference in the common parameter is determined to exceed the threshold (step 614, Figure 17);

- comparing the determined exothermic condition temperature with an exothermic condition (550 in Figure 17) expected from the catalyst at a time prior to the determined exothermic condition; and

- modifying the injected hydrocarbon in accordance with the last-mentioned comparison (steps 618 and 620; also see Figure 18 and line 10 of column 9 to line 3 of column 10) (Hirota et al. determine in advance a desired lower limit catalyst inlet temperature T1 and a desired upper limit catalyst outlet temperature T2 for the optimum reduction of NOx as a function of the degradation extent DR (Figure 17). For a non-deteriorated catalyst, T1 and T2 equal 450 and 550, respectively. If a detected temperature difference (Δt) across the catalyst is different from a predetermined temperature threshold (ΔTi), a degradation extent DR is calculated (step 612); and a set of desired temperature values T1 and T2 are determined based on the calculated DR (step 614). A hydrocarbon concentration H1 is also determined based on DR).

Re claims 8 and 11, in the system and method of Hirota et al., the common parameter is temperature and wherein the sensors are temperature sensors.

Response to Arguments

8. Applicant's arguments with respect to the references applied in the previous Office Action have been fully considered but they are not persuasive.

Re claims 1, 4, 12, and 20, in response to applicant's argument that Bagley et al. fail to detect an exothermic reaction across the catalyst (page 10 of Applicant's Amendment), the examiner respectfully disagrees.

Step (b) in claim 1 in Bagley et al. reads "applying sufficient energy to the electrically heated catalyst to attain the conversion temperature or ignition of the air/fuel mixture;". This

sentence means that Bagley et al. will turn off or reduce the electricity to the electrically heated

catalyst when the conversion temperature or ignition of the air/fuel mixture is observed or

detected (Also see steps (d) and (e) in claim 14). Figure 2 in Bagley et al. exhibits the exotherm

in a catalyst for three different fuels: methanol, ethanol, and hexane. Based on this figure,

Bagley et al. are able to determine that the conversion temperature for the catalyst they are

testing is at about 45°C for methanol, 50-60°C for ethanol, and about 160 to 170°C for hexane

(lines 35-37 of column 7). Bagley et al. further disclose that the conversion temperature of a

given fuel is a function of at least the aging of a catalyst (lines 31-34 of column 1). Thus, the

conversion temperatures of the three different fuels given above by Bagley et al. are not constant

values for any catalyst. They are at least a function of the deteriorated condition of the catalyst.

As illustrated in Figure 2, one of the more reliable ways to determine this conversion temperature

is provided by Bagley et al. By detecting an exotherm or a sharp temperature rise in the catalyst

when the catalyst reaches 45°C, Bagley et al. are able to determine that with methanol as a fuel,

the conversion temperature for a mixture of air and methanol for the specific catalyst they are

testing is at about 45°C. Thus, Bagley et al. clearly disclose the step in dispute.

Re claims 4-20, in response to applicant's argument that Hirota et al. fail to inject a hydrocarbon based on a measured temperature of the catalyst when an exothermic reaction is detected (pages 11 and 13 of Applicant's Amendment), the examiner again respectfully disagrees

As shown in Figure 14 and indicated on lines 28-33 of column 9, Hirota et al. detect a temperature difference or an exotherm ($\Delta t = t2 - t1$) indicating an exothermic reaction across the catalyst (step 608), wherein <u>t1</u> is a measured temperature of an inlet of the catalyst (6) and <u>t2</u> is a measured temperature of an output of the catalyst (emphasis added). Also in this step, the temperature of an output of the catalyst (t2) is measured in response to the detected exothermic reaction. Based on the detected exotherm Δt and the measured temperatures t1 and t2, Hirota et al. adjust and inject a hydrocarbon fuel into the catalyst (steps 618 and 620; also see Figure 18 and line 10 of column 9 to line 3 of column 10). Thus, Hirota et al. clearly disclose the claimed features of the pending application.

Conclusion

9. THIS ACTION IS MADE FINAL. Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Art Unit: 3748

Page 11

Communication

10. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Examiner Tu Nguyen whose telephone number is (571) 272-4862.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Mr. Thomas E. Denion, can be reached on (571) 272-4859. The fax phone number for the organization where this application or proceeding is assigned is (703) 872-9306.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

TMN

December 22, 2004

Tu M. Nguyen

Primary Examiner

Tu M. Nguyen

Art Unit 3748